Filing Date: October 31, 2006

## AMENDMENTS TO THE CLAIMS

 (Currently amended) An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of:

- (a) a nucleotide sequence that encodes a polypeptide conferring disease <u>fusarium</u> resistance to a plant, <u>or a full length complement of the nucleotide sequence</u>, <u>wherein the nucleotide sequence is selected from the sequence sharing at least-95% sequence identity</u> with the sequence set forth in SEQ ID NO: 1 or 3-or a complement thereof:
- (b) a nucleotide sequence that encodes a polypeptide conferring disease <u>fusarium</u> resistance to a plant and comprising an <u>the</u> amino acid sequence <del>that shares at least 95%</del> sequence identity with the sequence set forth in SEQ ID NO: 2 or 4, or a full length complement of the nucleotide sequence; and
- (c) a nucleotide sequence that encodes a polypeptide that confers disease <u>fusarium</u> resistance to a plant, <u>or a full length complement of the nucleotide sequence</u>, wherein the nucleotide sequence <u>hybridizes</u> hybridises to the sequence of (a) and (b), or to a <u>full length</u> complement of a nucleotide sequence selected from the group consisting of the sequence set forth in SEQ ID NO: 1 or 3 and a nucleotide sequence that encodes the amino acid sequence set forth in SEQ ID NO: 2 or 4, thereof, under high stringency conditions, wherein the conditions comprise hybridization at 65°C in 1% BSA, 1 mM EDTA, 0.5 M NaHPO<sub>4</sub> (pH 7.2), 7% SDS, and washing at 65°C in 0.2 X SSC, 0.1% SDS.
- (Original) A nucleic acid construct, comprising a polynucleotide according to claim 1 operably connected to a regulatory element, which is operable in the plant.
- (Original) A nucleic acid construct according to claim 2, wherein the construct is a vector.
- (Original) An isolated host cell containing a nucleic acid construct according to claim
  - 5. (Original) A host cell according to claim 4, wherein the host cell is a plant cell.

Filing Date: October 31, 2006

(Original) A host cell according to claim 5, wherein the plant cell has the nucleic acid construct incorporated into its nucleome.

 (Original) A host cell according to claim 5, wherein the plant cell has the nucleic acid construct stably incorporated into its genome.

8. (Original) A plant containing a cell comprising a nucleic acid construct according to

claim 2.

(Original) A plant according to claim 8, wherein the plant cell has the nucleic acid construct stably incorporated into its genome.

10. (Canceled)

11. (Canceled)

12. (Canceled)

13. (Canceled)

14. (Currently amended) A method for modulating disease resistance in a plant, the method comprising introducing a construct into the nucleome of the plant and regenerating a stably transformed plant, the construct comprising a regulatory element operably connected to a nucleotide sequence selected from the group consisting of: (a) a nucleotide sequence that encodes a polypeptide conferring disease fusarium resistance to a plant, wherein the nucleotide sequence is selected from the sequence set forth in SEQ ID NO:1 or 3 the sequence sharing at least 95% sequence identity with the sequence set forth in SEQ ID NO:1 or 3, or a complement thereof; (b) a nucleotide sequence that encodes a polypeptide conferring disease fusarium resistance to a plant and comprising an the amino acid sequence that shares at least 95% sequence identity with the sequence set forth in SEQ ID NO: 2 or 4; and (c) a nucleotide sequence that encodes a polypeptide that confers disease resistance to a plant, wherein the nucleotide sequence hybridizes hybridises to the full length complement sequence of (a) and or (b), or to a complement thereof; under high

Filing Date: October 31, 2006

stringency conditions, wherein the conditions comprise hybridization at 65°C in 1% BSA, 1 mM EDTA, 0.5 M NaHPO4 (pH 7.2), 7% SDS, and washing at 65°C in 0.2 X SSC, 0.1% SDS.

- 15. (Original) A method according to claim 14, wherein the construct is introduced into regenerable plant cells so as to yield transformed plant cells.
- 16. (Original) A method according to claim 15, wherein the transformed plant cells are used for regenerating a differentiated plant.
- 17. (Original) A method according to claim 15, wherein the regenerable cells are regenerable dicotyledonous plant cells.
- 18. (Original) A method according to claim 15, wherein the regenerable cells are regenerable monocotyledonous plant cells.
- (Currently amended) A method according to claim 15, wherein regenerable cells are regenerable graminaceous moneotyledonous plant cells.
- 20. (Original) A method according to claim 15, wherein regenerable cells are regenerable non-graminaceous monocotyledonous plant cells.
- (Original) A method according to claim 15, wherein regenerable cells are regenerable banana cells.
- 22. (Currently amended) A method according to claim 16, wherein the expression of the polynucleotide renders confers the differentiated transgenic plant with enhanced resistance to disease.
- (Original) A method according to claim 22, wherein disease is caused by a fungal pathogen.
- (Original) A method according to claim 22, wherein disease is caused by soil borne fungi.

Application No.: 10/573,372 Filing Date: October 31, 2006

 (Original) A method according to claim 22, wherein disease is caused by Fusarium species.

- 26. (Original) A method according to claim 16, wherein the nucleic acid construct is transmitted through a complete cycle of the differentiated transgenic plant to its progeny so that it is expressed by the progeny plants.
- 27. (Original) A method according to claim 26, wherein the progeny is selected from seed, plant parts, tissue, and progeny plants derived from the differentiated transgenic plant.
- 28. (Currently amended) A method of breeding a plant, the method comprising identifying a plant that is resistant to a pathogenie disease fusarium wilt by detecting expression in the plant of a polynucleotide; and transferring from the plant genetic material corresponding to the polynucleotide via crossing and backcrossing to another plant, wherein the polynucleotide comprises a nucleotide sequence that is selected from the group consisting of: (a) a nucleotide sequence that encodes a polypeptide conferring disease fusarium resistance to a plant, or a full length complement of the nucleotide sequence, wherein the nucleotide sequence is selected from the sequence set forth in SEQ ID NO: 1 or 3the sequence sharing at least 95% sequence identity with the sequence set forth in SEQ ID NO: 1 or 3, or a complement thereof; (b) a nucleotide sequence that encodes a polypeptide conferring disease fusarium resistance to a plant and comprising an the amino acid sequence that shares at least 95% sequence identity with the sequence set forth in SEQ ID NO: 2 or 4, or a full length complement of the nucleotide sequence; and (c) a nucleotide sequence that encodes a polypeptide that confers disease fusarium resistance to a plant, or a full length complement of the nucleotide sequence, wherein the nucleotide sequence hybridises hybridizes to a full length complement of a nucleotide sequence selected from the group consisting of the sequence set forth in SEQ ID NO: 1 or 3 and a nucleotide sequence that encodes the amino acid sequence set forth in SEQ ID NO: 2 or 4 the sequence of (a) or (b), or to a complement thereof, under high stringency conditions, wherein the conditions comprise hybridization at 65°C in 1% BSA, 1 mM EDTA, 0.5 M NaHPO<sub>4</sub> (pH 7.2), 7% SDS, and washing at 65°C in 0.2 X SSC, 0.1% SDS.

Filing Date: October 31, 2006

29. (Original) A method according to claim 28, wherein the other plant is susceptible to a pathogenic disease.

30. (Original) A method according to claim 29, wherein the disease is caused by a fungal

pathogen.

31. (Original) A method according to claim 29, wherein the disease is caused by a

Fusarium species.

32. (Original) A method according to claim 28, wherein the genetic material comprises

naturally-occurring DNA.

33. (Original) A method according to claim 28, comprising: (1) sexually crossing a plant

containing the genetic material with a plant from a pathogen susceptible taxon; (2) recovering

reproductive material from the progeny of the cross; and (3) growing plants with enhanced

resistance to the disease from the reproductive material.

34. (Canceled)

35. (Original) A method according to claim 33, further comprising repetitively: (a)

backcrossing the disease resistant progeny with disease susceptible plants from the susceptible

taxon; and (b) selecting for expression of a nucleic acid sequence corresponding to the polynucleotide or to marker gene associated with the polynucleotide among the progeny of the

-9-

backcross, until the desired characteristics of the susceptible taxon are present in the progeny.

36. (Canceled)

37. (Canceled)

38. (Canceled)

39. (Canceled)

40. (Canceled)

Application No.: 10/573,372 Filing Date: October 31, 2006

41. (Canceled)

42. (Canceled)

43. (Canceled)

44. (Canceled)

45. (Canceled)

46. (Canceled)

- 47. (Currently amended) An isolated polynucleotide comprising a nucleotide sequence encoding an amino acid sequence selecting selected from the group consisting of
- (i) an amino acid sequence which confers disease <u>fusarium</u> resistance to a plant, <u>wherein</u> the <u>amino acid sequence</u> is selected from the sequence set forth in SEQ ID NO: 2 or 4 and which shares at least 95% sequence identity with the sequence set forth in SEQ ID NO: 2 or 4;
- (ii) an amino acid sequence which confers disease <u>fusarium</u> resistance to a plant and which is encoded by [[a]] <u>the</u> nucleotide sequence that shares at least 95% sequence identity with the sequence set forth in SEQ ID NO: 1 or 3<sub>7</sub> or a complement thereof; <u>and</u>
- (iii) an amino acid sequence which confers disease <u>fusarium</u> resistance to a plant and which is encoded by a nucleotide sequence that <u>hybridises hybridizes</u> under high stringency conditions to <u>a full length complement of</u> the sequence set forth in SEQ ID NO: 1 or 3, or—a eemplement thereof wherein the conditions comprise hybridization at 65°C in 1% BSA, 1 mM EDTA, 0.5 M NaHPO<sub>4</sub> (pH 7.2), 7% SDS, and washing at 65°C in 0.2 X SSC, 0.1% SDS.

48. (Canceled)